



# Formycon AG The Biosimilar Experts

June 2025



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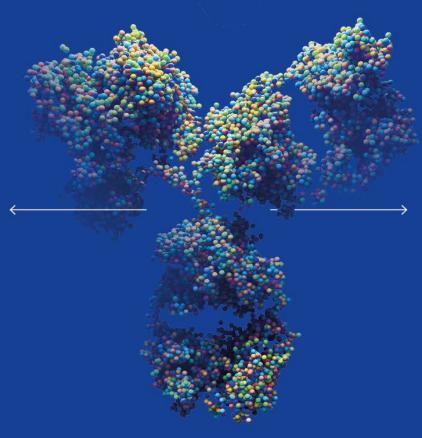
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### We are acting along a clear mission

### Biosimilars open up enormous opportunities

Contributing to ease the **financial strains** on the world's healthcare systems



Improving **patient access** to vital medicines

# **Skillset and mindset** are our key ingredients





Pure Play Biosimilar Company – established 2012 in Munich, Germany.

Business model contains Income from success payments and royalty streams.



**250 employees** from more than 30 different countries.

More than **80**% of Formycon's workforce is engaged in **R&D activities.** 



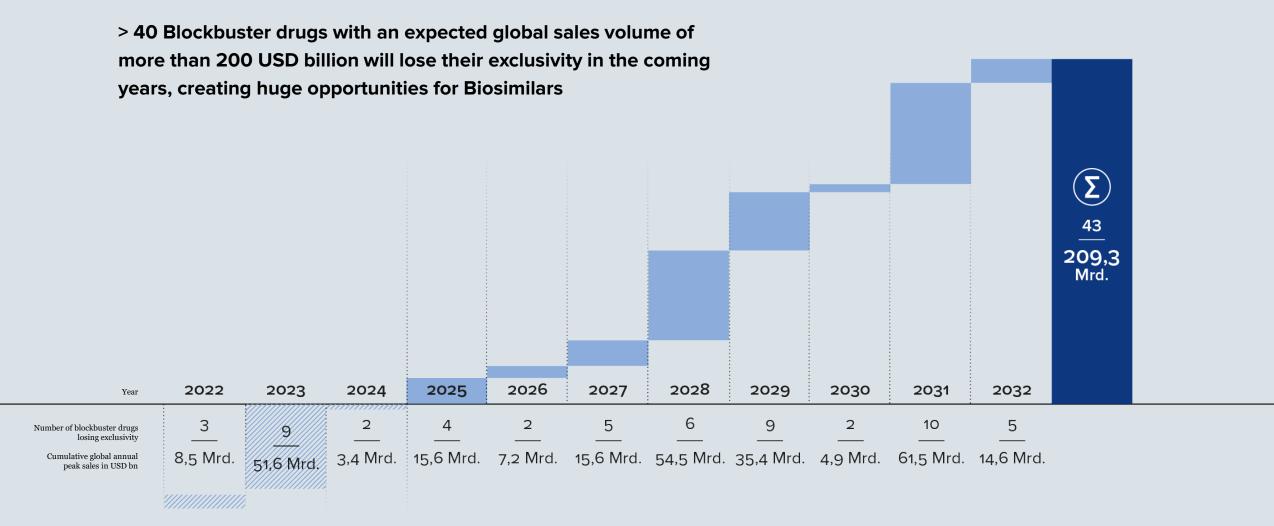
combining high professional expertise in biopharmaceutical development with agile mindset enables
Formycon to develop multiple Biosimilar projects in competitive timing and high quality.



Formycon's pipeline includes three approved biosimilars, two of which are already launched in key global markets, as well as four biosimilar candidates in development.

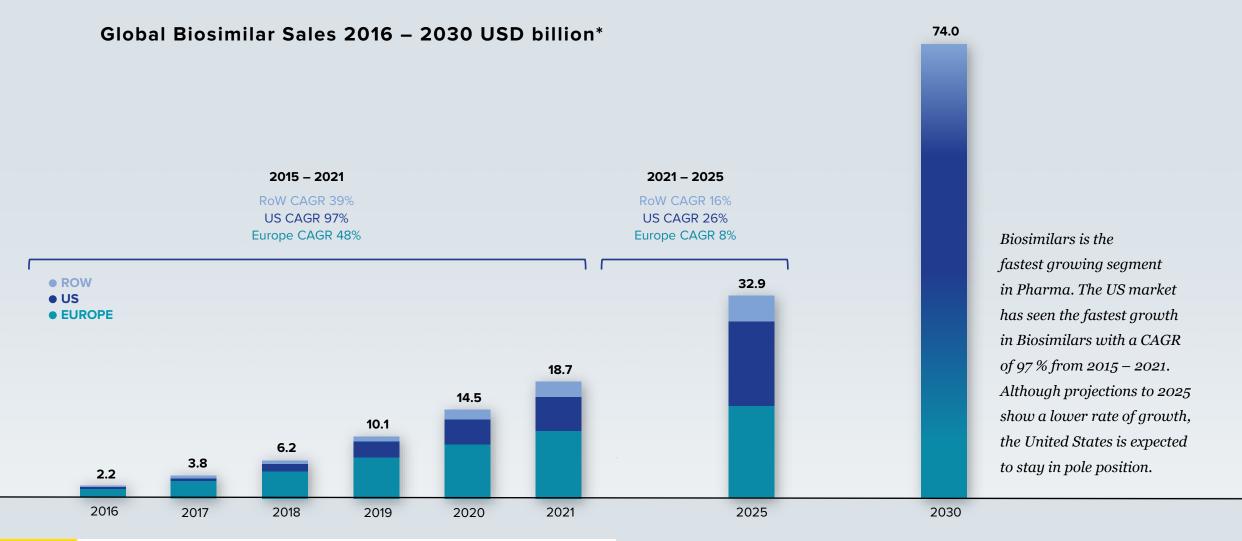


### **Huge Biosimilar target opportunities**





### The Biosimilar market develops very dynamically



# Laser focus on pipeline execution and commercial growth





### Maximizing our assets along a clear path

#### 2024

Important year with many operational milestones successfully achieved

#### 2025

Further transformation into a commercial company with two products on key global markets Achieving and growing sustainable profitability with maturing pipeline growth

Biosimilar Experts



### Strong maturing and growing pipeline

#### Diversified portfolio of commercial, late and mid stage programs



### Strong first half of 2025 – many important operational Milestones achieved





Approval of Stelara® Biosimilar FYB202/Otulfi® in Canada



Approval of Stelara® Biosimilar FYB202/Otulfi® in UK



Commercial Launch of Stelara® Biosimilar FYB202/Otulfi® in US, EU and Canada



Approval of
Eylea® Biosimilar
FYB203/AHZANTIVE®
in the EU and UK



Commercialization
Partnership with Valorum
for Eylea® Biosimilar
FYB203/AHZANTIVE®
in the US and Canada



Commercialization
Partnership with Teva
for Eylea® Biosimilar
FYB203/AHZANTIVE® in
major Parts of Europe



Commercialization
Partnership with Lotus
Pharmaceutical for
Eylea® Biosimilar
FYB203/AHZANTIVE® in
the APAC Region



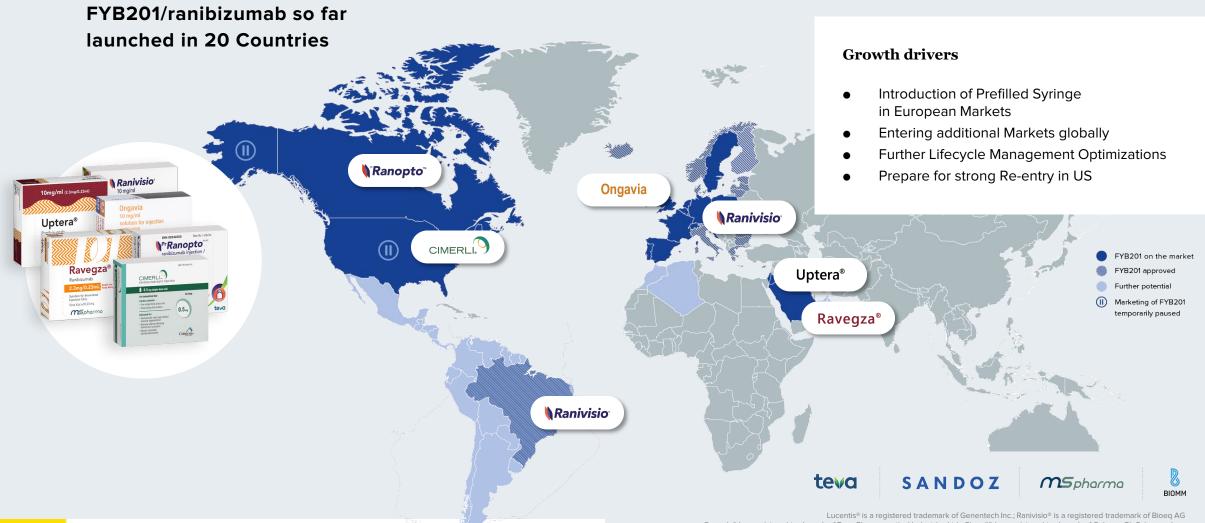
Tailored clinical Approach for Keytruda® Biosimilar Candidate FYB206 – without comparative efficacy (Phase-III) Study



Formycon included in the TecDAX Index of Deutsche Börse

# Lucentis® Biosimilar FYB201 – Strong Presence across the World





### Stelara® Biosimilar FYB202 – first Patients treated with Otulfi®





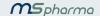
#### **Growth drivers**

- Build Momentum in US, EU and CA
- Add. Launches globally, e.g. UK
- Further LCM Optimizations

#### **Formycon Income Position**

- 2023 & 2024: Milestone payments in total of € 60m received
- 2024: One time revenue of approx. € 10m from selling "remaining development materials"
- From 2025 onwards: Post-commercialization value shared approximately equally by Formycon and Fresenius Kabi.

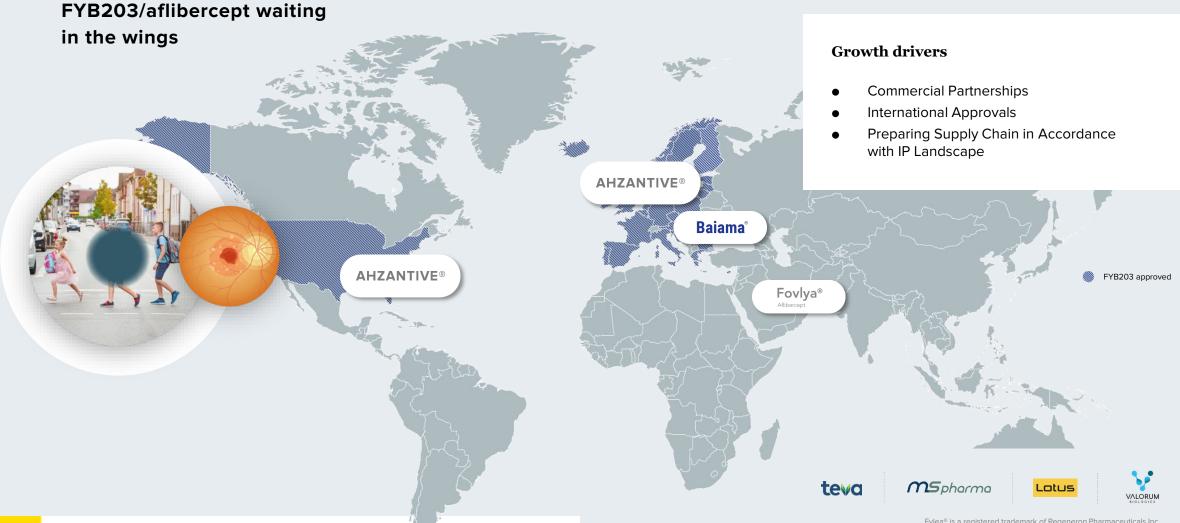






# Eylea® Biosimilar FYB203 – approved in US, EU and UK





# FYB206 – Keytruda® Biosimilar Candidate in the leading group





#### **Targeted Reference Indications**

Immuno-oncology: Melanoma (black skin cancer), non-small cell Lung Cancer, classical Hodgkin's Lymphoma and other Tumor Diseases

#### **Target Market 2024**

USD 29.5 billion

#### **Project Rights**

100% of project and commercialization rights

#### **Achievements and next important Milestones**

- Clinical Phase I trial "Dahlia" started in June 2024 investigating the PK equivalence as part of a preventive therapy for patients who have had a malignant melanoma (black skin cancer) completely surgically removed
- After in-depth scientific dialogue and in consultation with the FDA, Formycon has come to the conclusion that the Phase III clinical trial initiated in July 2024 is no longer necessary for the development and approval of FYB206 in the U.S. In February 2025, we therefore announced that we will discontinue the Phase III trial.
- Concluding regional or global commercialization partnerships





report/keytruda-market/
Keytruda® is a registered trademark
of Merck Sharp & Dohme LLC

### Outlook for the second half of 2025 – further significant commercial and operational milestones in sight





Approval of Lucentis® Biosimilar FYB201/Ranivisio® in LATAM



Approval and Launch of prefilled Syringe for Lucentis® Biosimilar FYB201



Commercial Launch of Stelara® Biosimilar FYB202/Otulfi® in the UK



Commercialization
Partnerships for Eylea®
Biosimilar
FYB203/AHZANTIVE®
in further Regions



Commercialization
Partnerships for Keytruda®
Biosimilar Candidate
FYB206



Disclosure of Immunology Biosimilar Candidate FYB208



... and many more important Milestones to come in the Course of 2025



#### 2025 outlook – Guidance confirmed

Guidance 2025	Revenue	EBITDA	Adjusted EBITDA	Working Capital
	<b>55</b> to <b>65</b>	-20 to -10	-20 to -10	<b>25</b> to <b>35</b>
	€ million	€ million	€ million	€ million
Key financial Figures Q1 2025	Revenue	EBITDA	Adjusted EBITDA	Working Capital
	5.3	-13.2	-11.8	29.4
	€ million	€ million	€ million	€ million
YE 2024		_	_	
	Revenue ———	EBITDA	Adjusted EBITDA	Working Capital
	69.7	-13.7	-1.6	55.1

€ million

€ million

#### **Guidance 2025**

#### Revenue:

€ million

• 1Q revenue as expected

#### **EBITDA:**

For Full Year expected on guidance

#### **Adjusted EBITDA**

- At Equity result above expectations in 1Q
- Expected to reverse during the year

#### **Working Capital:**

As expected

#### Liquidity

- End of Q1 25 total Cash reserves amounted to € 32.9m
- Plus undrawn €48m shareholder loan available

#### **Stable Guidance**

- Overall numbers are on track for Q1 2025
- Guidance 2025 confirmed

€ million

# Formycon – stable Anchor Investors and increased Liquidity



- Market Segment: Frankfurt Stock Exchange Regulated Market (Prime Standard)
- Uplisted to Prime Standard on Nov. 12, 2024,
   part of the SDAX since Dec. 23, 2024,
   joined the TecDAX on Jan. 13, 2025,
  - more international Investors
  - higher Liquidity
  - better Transparency
- Registered capital: € 17,664,427
   Shares outstanding: 17,664,427 (w/o par value)
- Market price / Market capitalization: ~ € 400 million
- Member of Indices: SDAX, TecDax, MSCI Europe Small Cap,
   MSCI EAFE IMI, MSCI Germany Small Cap

#### **Shareholder Structure**

- 24.04 % Santo Holding (Deutschland) GmbH
- 13.25 % Wpart GmbH, Wen.Co Invest GmbH, Peter Wendeln
- 9.08 % Gedeon Richter
- 6.04 % Active Ownership
- **5.10** % Detlef & Ursula Spruth
- **3.28** % Stefan R.
- **39.21** % Free Float\*\*



#### Research coverage:

– Berenberg	Виу	<ul> <li>Metzler Capital Markets</li> </ul>	Buy
<ul><li>First Berlin</li></ul>	Виу	– M. M. Warburg	Виу
<ul> <li>Hauck Aufhäuser</li> </ul>	Виу	<ul> <li>mwb Research</li> </ul>	Buy
<ul> <li>HC Wainwright</li> </ul>	Виу	<ul><li>Oddo BHF</li></ul>	Neutral
– Jefferies	Виу	<ul> <li>Royal Bank of Canada</li> </ul>	Виу
<ul> <li>Kepler Cheuvreux</li> </ul>	Виу		



Herzlich willkommen im Prime Standard Formycon AG Kürzel: FYB WKN: AIEWVY Sektor: Pharma & Healthcare Subsektor: Biotechnolosie





# Fully focused Pure-Play Biosimilar Company





WE HAVE all ingredients to successfully develop and commercialize a growing pipeline



WE ACT in a highly attractive market



WE CREATED
a strong Platform with
track record



WE ARE entering the next stage of the Formycon Growth Story



### I AM HAPPY TO ANSWER YOUR QUESTIONS

www.formycon.com









### Formycon AG



#### **Formycon AG**

Fraunhoferstr. 15 82152 Planegg-Martinsried Germany

+ 49 89 864 667 100 info@formycon.com

www.formycon.com





